- 10. The method of Claim 9, wherein the cell division inhibitor is selected from the group consisting of chloramphenicol, a protein synthesis inhibitor, an organic compound having β-lactamase inhibiting ability, nalidixic acid, promidic acid, pipemidic acid, oxolinaic acid, ofloxacin and enoxacin.
- 11. The method of Claim 10, wherein the protein synthesis inhibitor is selected from the group consisting of tetracycline, puromycin and erythromycin.
- 12. The method of Claim 10, wherein the organic compound having β -lactamase inhibiting ability is thienamycin.
 - 13. The method of Claim 10, wherein the cell division inhibitor is nalidixic acid.
- 14. The method of Claim 9, wherein the concentration of the cell division inhibitor in the culture medium is 0.01 to 5 mM.
 - 15. The method of Claim 9, wherein the bacteria are Acetobacter.
- 16. The method of Claim 15, wherein the bacteria are *Acetobacter pasteurianus* FERM BP-4176.
- 17. The method of Claim 9, further comprising recovering the bacterial cellulose produced in the culture medium.
 - 18. The method of Claim 10, wherein the cell division inhibitor is chloramphenicol.
- 19. A bacterial cellulose which is produced by culturing cellulose-producing bacteria which produce the bacterial cellulose extracellularly in a culture medium containing a cell division inhibitor.--